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ACE2 links amino acid malnutrition to microbial ecology and intestinal inflammation

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Abstract: Malnutrition affects up to one billion people in the world and is a major cause of mortality. In many cases, malnutrition is associated with diarrhoea and intestinal inflammation, further contributing to morbidity and death. The mechanisms by which unbalanced dietary nutrients affect intestinal homeostasis are largely unknown. Here we report that deficiency in murine angiotensin I converting enzyme (peptidyl-dipeptidase A) 2 (Ace2), which encodes a key regulatory enzyme of the renin-angiotensin system (RAS), results in highly increased susceptibility to intestinal inflammation induced by epithelial damage. The RAS is known to be involved in acute lung failure, cardiovascular functions and SARS infections. Mechanistically, ACE2 has a RAS-independent function, regulating intestinal amino acid homeostasis, expression of antimicrobial peptides, and the ecology of the gut microbiome. Transplantation of the altered microbiota from Ace2 mutant mice into germ-free wild-type hosts was able to transmit the increased propensity to develop severe colitis. ACE2-dependent changes in epithelial immunity and the gut microbiota can be directly regulated by the dietary amino acid tryptophan. Our results identify ACE2 as a key regulator of dietary amino acid homeostasis, innate immunity, gut microbial ecology, and transmissible susceptibility to colitis. These results provide a molecular explanation for how amino acid malnutrition can cause intestinal inflammation and diarrhoea.

DOI: <https://doi.org/10.1038/nature11228>

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ZORA URL: <https://doi.org/10.5167/uzh-65968>
Journal Article

Originally published at:

Hashimoto, Tatsuo; Perlot, Thomas; Rehman, Ateequr; Trichereau, Jean; Ishiguro, Hiroaki; Paolino, Magdalena; Sigl, Verena; Hanada, Toshikatsu; Hanada, Reiko; Lipinski, Simone; Wild, Birgit; Camargo, Simone M R; Singer, Dustin; Richter, Andreas; Kuba, Keiji; Fukamizu, Akiyoshi; Schreiber, Stefan; Clevers, Hans; Verrey, Francois; Rosenstiel, Philip; Penninger, Josef M (2012). ACE2 links amino acid malnutrition to microbial ecology and intestinal inflammation. *Nature*, 487(7408):477-481.

DOI: <https://doi.org/10.1038/nature11228>

ACE2 links amino acid malnutrition to microbial ecology and intestinal inflammation

Tatsuo Hashimoto^{1,2*}, Thomas Perlot^{1*}, Ateequr Rehman^{3†}, Jean Trichereau¹, Hiroaki Ishiguro², Magdalena Paolino¹, Verena Sigl¹, Toshikatsu Hanada¹, Reiko Hanada¹, Simone Lipinski³, Birgit Wild⁴, Simone M. R. Camargo⁵, Dustin Singer⁵, Andreas Richter⁴, Keiji Kuba⁶, Akiyoshi Fukamizu⁷, Stefan Schreiber², Hans Clevers⁸, Francois Verrey⁵, Philip Rosenstiel³, and Josef M. Penninger¹

¹ IMBA, Institute of Molecular Biotechnology of the Austrian Academy of Sciences, 1030 Vienna Austria.

² The Department of Medical Science and Cardiorenal Medicine, Yokohama City University Graduate School of Medicine and School of Medicine, Yokohama, Japan

³ Institute of Clinical Molecular Biology, University Kiel, Schittenhelmstr.12, 24105, Germany

⁴ Department of Chemical Ecology & Ecosystem Research, Center of Ecology, University of Vienna, Althanstrasse. 14, A-1090 Vienna, Austria.

⁵ Institute of Physiology and Center for Integrative Human Physiology, University of Zurich, Winterthurerstrasse 190, 8057 Zurich, Switzerland

⁶ Department of Biological Informatics and Experimental Therapeutics, Akita University Graduate School of Medicine, 1-1-1 Hondo, Akita, 010-8543, Japan.

⁷ Graduate School of Life and Environmental Sciences, University of Tsukuba, Tsukuba, Ibaraki 305-8577, Japan

⁸ Hubrecht Institute, Royal Netherlands Academy of Arts and Sciences and University Medical Center Utrecht, Utrecht, The Netherlands

† Present address: Department of Environmental Health Sciences, University Medical Center, Breisacher Str. 115b, D-79106 Freiburg, Germany

* These authors contributed equally to this work

Malnutrition affects hundreds of millions of people in the world and is a major cause of mortality^{1,2}. In many cases malnutrition is associated with diarrhea and intestinal inflammation further contributing to morbidity and death². The mechanisms by which unbalanced dietary nutrients affect intestinal homeostasis are largely unknown. Here we report that deficiency in murine Angiotensin-converting enzyme 2 (*Ace2*), encoding a key regulatory enzyme of the renin-angiotensin system (RAS) involved in acute lung failure³, cardiovascular functions⁴, and SARS infections⁵, results in highly increased susceptibility to intestinal inflammation induced by epithelial damage. Mechanistically, ACE2 exhibits a RAS-independent function regulating intestinal amino acid homeostasis, expression of antimicrobial peptides, and the ecology of the gut microbiome. Transplantation of the altered microbiota from *Ace2* mutant mice into germfree wild type hosts was able to transmit the increased propensity to develop severe colitis. ACE2-dependent changes in epithelial immunity and the gut microbiota can be directly regulated by the dietary amino acid tryptophan. Our results identify ACE2 as a key regulator of dietary amino acid homeostasis, innate immunity, gut microbial ecology, and transmissible susceptibility to colitis. These results provide a molecular explanation how amino acid malnutrition can cause intestinal inflammation and diarrhea.

The renin-angiotensin system (RAS) has now been studied for more than a century. The RAS is regulated by the opposing actions of two key carboxypeptidases, angiotensin converting enzyme (ACE) and ACE2^{ref.4,6,7}. In addition to its catalytic activity, we and others have recently found that ACE2 associates with the neutral amino acid transporter B⁰AT1 and that ACE2 is required for expression of this transporter on the luminal surface of intestinal epithelial cells^{8,9}. Variants in the B⁰AT1 (*SLC6A19*) gene have been identified as a cause of Hartnup's disease¹⁰, a rare autosomal recessive disorder associated with pellagra-like symptoms including diarrhea that manifests under conditions of malnutrition¹¹. However, the *in vivo* function of ACE2 in the gut epithelium remained to be investigated.

Morphological and ultra-structural analysis of the small and large intestine of *Ace2* knock-out mice did not reveal any alterations. Intriguingly, when we challenged *Ace2* deficient mice with dextran sodium sulfate (DSS), an irritant that disrupts the intestinal epithelial barrier and results in colitis¹², a profoundly increased inflammatory reaction was observable compared to wild type littermates. We detected an enhanced infiltration of inflammatory cells, significant shortening of the colon length, increased intestinal bleeding (hemoccult), crypt damage, inflammation (redundant?), weight loss, and severe diarrhea resulting in an overall increased disease activity¹³ (Fig. 1a-c, Supplementary Fig. 1). Similar results were obtained using Trinitrobenzene sulphonic acid (TNBS) -induced colitis (Supplementary Fig. 2). Thus, genetic inactivation of the key RAS enzyme ACE2 results in severe colitis following intestinal injury.

To test whether the severe colitis phenotype is due to the catalytic activity of ACE2, we

treated mice with recombinant soluble ACE2 (rsACE2). We and others have shown that rsACE2 rescues virtually all previously reported *in vivo* ACE2 functions within the RAS system^{3,14}. As it lacks the transmembrane domain, soluble ACE2 cannot associate with the neutral amino acid transporter B⁰AT1 and thus cannot stabilize its cell-surface localization⁸. Treatment of mice with rsACE2 did not rescue the severe DSS-induced colitis phenotype of *Ace2* mutant mice (Supplementary Fig. 3a-h). One *in vivo* consequence of *Ace2* deletion within the RAS is accumulation of angiotensin II (AngII), which then primarily acts on the G-protein coupled AT1a receptor (*Agtr1a*)^{ref.3}. We indeed observed increased AngII levels in the colon of DSS-treated *Ace2* mutant mice that were reduced to background levels after treatment with rsACE2 (Supplementary Fig. 3i). To rule out that locally increased AngII might cause severe colitis, we generated *Agtr1a*^{-/-}*Ace2*^{-/-} mice. Deletion of *Agtr1* did not rescue the severe DSS-induced colitis (Supplementary Fig. 4), indicating that these effects are independent of the classical RAS system. Whether ACE inhibition might alter the phenotype of *Agtr1a*^{-/-}*Ace2*^{-/-} mice needs to be further examined. In addition to cleaving AngII, ACE2 exhibits catalytic activity towards a second peptide system, Apelin¹⁵. However, DSS-induced colitis was not altered in mice carrying genetic mutations in *Apelin* (Supplementary Fig. 5) or its receptor *Apj* (Supplementary Fig. 6). Thus, the catalytic activity of ACE2, essential for its function in the RAS and Apelin cleavage, has no overt role in DSS-induced intestinal inflammation.

It had been reported that the RAS can control immune functions¹⁶. However, in unchallenged *Ace2* mutant mice, we did not observe any apparent differences in immune cell populations of the colon and small intestine (insert: “not shown”).

TLR1-9 mRNA levels in jejunum, ileum, and colon were largely similar in *Ace2* deficient and wild type mice, with the exception of TLR5 and TLR9 in the colon (Supplementary Fig. 7). Transplantation of bone marrow from *Ace2* mutant mice (KO) into wild type (WT) mice (KO → WT) did not result in any obvious effect on DSS-induced colitis as compared to WT → WT transplant recipients. By contrast, reconstitution of *Ace2* mutant mice with WT bone-marrow (WT → KO) led to an increased severity of DSS-induced colitis that was indistinguishable from the KO → KO control group (Supplementary Fig. 8). Thus, ACE2 exerts its role on intestinal inflammation in non-haematopoietic cells.

ACE2 expression was primarily mapped to the luminal surface of differentiated small intestinal epithelial cells, whereas lower ACE2 expression is observed in crypt cells and the colon (Supplementary Fig. 9a,b & 10a). *Ace2* inactivation had no overt effect on basal proliferation of small intestinal or colon crypt cells (Supplementary Fig. 10a,b) nor did it affect apoptosis rates of intestinal epithelial cells (Supplementary Fig. 10a,c). ACE levels were slightly, albeit not significantly, increased in the jejunum of *Ace2*^{-/-} mice (Supplementary Fig. 9c,d). As reported previously⁸, protein expression of the neutral amino acid transporter B⁰AT1 was absent in the small intestine of *Ace2* mutant mice, whereas mRNA expression was not affected (Supplementary Figs. 9e,f & 11a). Due to the lack of intestinal B⁰AT1 protein expression, serum levels of the neutral amino acids valine (Val), threonine (Thr), and tyrosine (Tyr), and the essential amino acid tryptophan (Trp) were markedly reduced in *Ace2*^{-/-} mice (Supplementary Fig. 11b). The marked reduction in tryptophan correlates with human data, as patients suffering from Hartnup's disease caused by *B⁰AT1* mutations or malnutrition such as in anorexia

nervosa exhibit reduced tryptophan serum levels^{17,18}.

ACE2 is a chimeric protein that emerged from the duplication and fusion of two genes: homology with ACE at the catalytic domain and with Collectrin (Tmem27) in the membrane proximal domain¹⁹. Inactivation of *Collectrin* in mice results in a near complete downregulation of apical amino acid transporters such as B⁰AT1 in the kidney thereby regulating renal amino acid re-absorption¹⁹. By contrast, ACE2 associates with B⁰AT1 on the luminal surface of intestinal epithelial cells, i.e. ACE2/B⁰AT1 regulate the uptake of neutral amino acids in the intestine where Collectrin is not expressed (Supplementary Fig. 11a,c & 12). In both *Collectrin* and *Ace2* mutant mice, tryptophan is reduced in the serum (Supplementary Fig. 11b and ref. 19). To test whether impaired re-absorption of amino acids in the kidney might also result in altered susceptibility to colitis, we challenged *Collectrin*^{-/-} mice with DSS. However, genetic ablation of *Collectrin* did not affect DSS-induced colitis (Supplementary Fig. 13). Although genetic inactivation of both *Ace2* or *Collectrin* result in a similar deregulation of amino acids in the serum, functionally only the loss of ACE2 affects the susceptibility to intestinal inflammation.

We, therefore, speculated that the function of ACE2 in intestinal uptake of dietary amino acids might underlie the observed phenotype. To test this hypothesis, we fed *Ace2* mutant and control mice a protein free-diet (PFD) to eliminate dietary amino acids and challenged these mice with DSS at a dose (1%) that triggers only very mild colitis. Basal weight loss was comparable among control and *Ace2* mutant mice during the period of the PFD (Supplementary Fig. 14a). Moreover, average food intake did not

show a significant difference between *Ace2* deficient and control mice fed PFD or normal chow (Supplementary Fig. 14b). Importantly, PFD markedly worsened DSS-triggered colitis in wild type mice to levels seen in *Ace2* mutants (Fig. 1d-f; Supplementary Fig. 14c-g). Thus, protein malnutrition alters the severity of DSS-induced intestinal inflammation.

The general lack of dietary protein might induce broad effects on several organ systems. We, therefore, set out to define whether a specific amino acid might be responsible for the severe inflammatory phenotype. Since serum tryptophan was markedly decreased in *Ace2* mutant mice, we focused on this essential amino acid. Tryptophan is required for the *in vivo* generation of nicotinamide (vitamin B3/niacin)¹⁷ and insufficient niacin or tryptophan in the diet are the cause of pellagra, a disease still endemic in many countries with protein malnutrition²⁰. More than 90% of pellagra patients develop colitis¹¹ and for nearly 80 years nicotinamide (NAM) has been used as treatment for pellagra. In *Ace2* mutant mice, NAM almost completely alleviated the severe colitis and diarrhea (Fig. 2a-c; Supplementary Fig. 15). To demonstrate that the severe colitis in *Ace2* deficient mice is due to impaired tryptophan uptake, we provided tryptophan in form of a Gly-Trp dipeptide to bypass the loss of the single amino acid transporter B⁰AT1. Dietary dipeptidic tryptophan (Trp+) restored serum tryptophan levels (Supplementary Fig. 16a) and rescued the enhanced DSS-susceptibility of *Ace2*^{-/-} mice (Fig. 2d-f; Supplementary Fig. 16b-d). Tryptophan free (Trp-) diet resulted in a dramatic increase in susceptibility to DSS-induced inflammation (Supplementary Fig. 17a-h). In addition, Trp- diet markedly worsened weight loss in wild type mice challenged with TNBS (Supplementary Fig. 17i-l). Our results show that deficiency of *Ace2* causes a critical

impairment of local tryptophan homeostasis which alters the susceptibility to intestinal inflammation.

It is conceivable that an altered amino acid availability in the epithelium could lead to decreased regenerative responses and repair mechanisms after epithelial injury^{21,22}. However, proliferation rates and cell death of small intestinal and colon epithelial cells were comparable between control and *Ace2* mutant mice following DSS challenge (Supplementary Fig. 18). However, expression of multiple antimicrobial peptides was markedly reduced in isolated gut ileal epithelial cells from unchallenged *Ace2* mutant mice (Fig. 3a; Supplementary Fig. 19a-c). Tryptophan-free (Trp-) diet of wild type mice also resulted in a significant downregulation of antimicrobial peptides (Fig. 3b). Similar downregulation of antimicrobial peptides was observed upon feeding wild type mice a protein-free diet (Supplementary Fig. 20a). By contrast, dietary NAM (Supplementary Fig. 20b-g) and dipeptidic tryptophan (Trp+) (Fig. 3c and Supplementary Fig. 20h,i) triggered induction of antimicrobial peptides in intestinal epithelial cells from *Ace2* mutant and wild type mice. Thus, dietary tryptophan controls expression of small intestinal antimicrobial peptides.

Amino acids and nicotinamide can activate mTOR, involved in cell proliferation, survival, protein synthesis and transcription²³. In epithelial intestinal cells from unchallenged *Ace2* mutant mice we observed markedly impaired p70S6 kinase activity (Supplementary Fig. 21a) as well as reduced S6 phosphorylation (Fig. 3d and Supplementary Fig. 21b), indicative of reduced mTOR activity. A similar reduction of mTOR activity was recently reported in mice lacking B⁰AT1^{ref.24}. Administration of

dipeptidic tryptophan (Trp+) resulted in increased mTOR activity in the small intestine of *Ace2* mutant mice (Fig. 3e). *In vivo* inhibition of mTOR with rapamycin (RAPA) in wild type mice resulted in a significant downregulation of antimicrobial peptide expression, an effect that could not be rescued by NAM (Supplementary Fig. 22a). Moreover, administration of RAPA before the first challenge with DSS increased the severity of colitis (Fig. 3f; Supplementary Fig. 22b-h). Of note, although acute mTOR inhibition may have beneficial effects in murine colitis models²⁵, mTOR blockade has not been proven successful in human clinical trials in inflammatory bowel disease (IBD)²⁶. We could delete this sentence if still too long? Thus, at the molecular level, nicotinamide and dietary tryptophan appear to exert their effects on intestinal antimicrobial peptides and colitis via the mTOR pathway.

How does ACE2 regulated uptake of tryptophan in the small intestine affect DSS-induced inflammation in the colon? It has been shown that α -defensins secreted into the ileum lumen persist in a functional state throughout the entire gut²⁷ and that alterations in antimicrobial peptides can affect the ecology of the small and large bowel microbiota²⁸. We therefore performed deep profiling of the intestinal microbiome of *Ace2* mutant and wild type littermates using 16s rDNA fingerprinting. The luminal ileocecal microbiome of *Ace2* mutant mice was markedly altered as shown by weighted (Bray-Curtis) (Fig. 4a) and unweighted (Unifrac) (Fig. 4b) analyses. Distinct operational taxonomical units (OTUs) are overrepresented in *Ace2* mutant mice (Fig. 4c; Supplementary Tables 1 & 2). RAPA treatment resulted in a distinct alteration of the ileocecal gut microbiome in wild type animals which was still closer related to wild type than to *Ace2*^{-/-} animals (Supplementary Fig. 23a-c; Supplementary Tables 3 & 4); RAPA

acts on many different cell types that in addition to the changes in intestinal epithelial cells could alter microbiota composition²⁹. Importantly, both dipeptidic tryptophan (Trp+) and NAM treatment reverted the composition of the intestinal microbiota of *Ace2* mutant mice to be more similar to wild type littermates (Fig. 4a-c, Supplementary Fig. 24a-c, Supplementary Tables 1, 2, 5 & 6). Thus, de-regulation of tryptophan amino acid homeostasis in *Ace2* mutant mice alters the intestinal microbiome.

Consistent with an alteration in the intestinal microbiome, antibiotic treatment of *Ace2* mutant mice alleviated the severe colitis (Supplementary Fig. 25). We finally performed gut microbiome transplants from control and *Ace2* mutant mice into germfree hosts followed by DSS challenge. Transplantation of the ileocecal gut microbiota from *Ace2* mutant mice transmitted the inflammatory phenotype to wild type hosts as compared to mice that were colonized with the microbiota of *Ace2*^{+/-} mice (Fig. 4d,e; Supplementary Fig. 26a-f). Expression of antimicrobial peptides in wild type mice did not change significantly upon transplantation of *Ace2*^{+/-} or *Ace2*^{-/-} microbiota within the timeframe of analysis (Supplementary Fig. 26g).

We propose the following scenario how malnutrition and tryptophan deficiency can cause diarrhea and colitis. Dietary tryptophan is primarily absorbed via the B⁰AT1/ACE2 transport pathway on the luminal surface of small intestinal epithelial cells. This results in the activation of mTOR either directly through nutrient sensing and/or through the tryptophan-nicotinamide pathway. mTOR then regulates expression of antimicrobial peptides which affect the intestinal composition of the gut microbiota. Whether pathways other than mTOR and antimicrobial peptides also contribute to the

altered microbiome needs to be explored. Our results further indicate that under conditions of intestinal injury, such an altered microbiome contributes to the severity of colitis. Importantly, our data indicate that the essential amino acid tryptophan and its metabolite nicotinamide are key regulators of gut microbiota and propensity to inflammation.

Malnutrition is a major global health burden affecting up to one billion people in the world and its consequences such as colitis and diarrhea are often fatal^{1,2}. In addition, chronic inflammatory conditions of the intestine, cachexia, anorexia nervosa, or amino acid mal-absorption constitute severe clinical problems³⁰. Our results provide novel mechanistic insights how protein malnutrition can lead to colitis and diarrhea. Our data also identify a molecular crosstalk between the RAS system and intestinal amino acid homeostasis via ACE2 and provide a direct link of dietary amino acid metabolism to innate immunity, the composition of the intestinal microbiota, and susceptibility to colitis.

METHODS SUMMARY

Colitis models. *Ace2*, *Collectrin*, *Agtr1*, *Apj*, and *Apelin* mutant mice have been described. Mice were given protein-free and tryptophan-free diet, tryptophan di-peptides, NAM, rapamycin, or were treated with catalytically active murine ACE2 protein and challenged with DSS or TNBS. Bone marrow transplantations were performed in lethally irradiated *Ace2* mutant and wild type mice. For microbiota transplantation, cecal and ileum extracts from *Ace2*^{+/y} or *Ace2*^{-/y} mice were transplanted to germ free hosts. Severity of colitis was monitored as described¹³. All experiments were performed in accordance with institutional guidelines.

Expression analyses. Small and large intestinal epithelial cells and mucosal cells were purified and gene expression assessed using qRT-PCR. Protein expression was determined by Western blot and localized using immunohistochemistry. Immune cell populations were determined by FACS.

Microbiota sequencing. Genomic DNA was extracted from the terminal ileum, amplified by PCR, and sequenced using Roche 454 titanium chemistry. All sequence reads were filtered for quality and length using PANGEA. Sequences were clustered using average neighbour algorithm into species level operational taxonomical units (OTUs). Distance matrices between samples were generated based on weighted (Bray-Curtis similarity) and non-weighted (unweighted UniFrac) algorithms and shown as principal coordinate analysis (PCoA). Spearman correlations and similarity percentage analysis (SIMPER) were performed to determine significant differences among bacterial communities among genotype and diet groups.

Full Methods and any associated references are available in the online version of the paper at www.nature.com/nature.

- ¹ Khan, Y. & Bhutta, Z.A., Nutritional deficiencies in the developing world: current status and opportunities for intervention. *Pediatr Clin North Am* **57**, 1409-1441 (2010).
- ² Weisstaub, G. & Araya, M., Acute malnutrition in Latin America: the challenge of ending avoidable deaths. *J Pediatr Gastroenterol Nutr* **47**, S10-14 (2008).
- ³ Imai, Y. *et al.*, Angiotensin-converting enzyme 2 protects from severe acute lung failure. *Nature* **436**, 112-116 (2005).
- ⁴ Crackower, M.A. *et al.*, Angiotensin-converting enzyme 2 is an essential regulator of heart function. *Nature* **417**, 822-828 (2002).
- ⁵ Kuba, K. *et al.*, A crucial role of angiotensin converting enzyme 2 (ACE2) in SARS coronavirus-induced lung injury. *Nat Med* **11**, 875-879 (2005).
- ⁶ Donoghue, M. *et al.*, A novel angiotensin-converting enzyme-related carboxypeptidase (ACE2) converts angiotensin I to angiotensin 1-9. *Circ Res* **87**, E1-9 (2000).
- ⁷ Tipnis, S.R. *et al.*, A human homolog of angiotensin-converting enzyme. Cloning and functional expression as a captopril-insensitive carboxypeptidase. *J Biol Chem* **275**, 33238-33243 (2000).
- ⁸ Camargo, S.M. *et al.*, Tissue-specific amino acid transporter partners ACE2 and collectrin differentially interact with hartnup mutations. *Gastroenterology* **136**, 872-882 (2009).
- ⁹ Kowalczyk, S. *et al.*, A protein complex in the brush-border membrane explains a Hartnup disorder allele. *FASEB J* **22**, 2880-2887 (2008).

- 10 Kleta, R. *et al.*, Mutations in SLC6A19, encoding B0AT1, cause Hartnup disorder. *Nat Genet* **36**, 999-1002 (2004).
- 11 Segal, I. *et al.*, Rectal manifestations of pellagra. *Int J Colorectal Dis* **1**, 238-243 (1986).
- 12 Okayasu, I. *et al.*, A novel method in the induction of reliable experimental acute and chronic ulcerative colitis in mice. *Gastroenterology* **98**, 694-702 (1990).
- 13 Cooper, H.S., Murthy, S.N., Shah, R.S., & Sedergran, D.J., Clinicopathologic study of dextran sulfate sodium experimental murine colitis. *Lab Invest* **69**, 238-249 (1993).
- 14 Osterreicher, C.H. *et al.*, Angiotensin-converting-enzyme 2 inhibits liver fibrosis in mice. *Hepatology* **50**, 929-938 (2009).
- 15 Vickers, C. *et al.*, Hydrolysis of biological peptides by human angiotensin-converting enzyme-related carboxypeptidase. *J Biol Chem* **277**, 14838-14843 (2002).
- 16 Swirski, F.K. *et al.*, Identification of splenic reservoir monocytes and their deployment to inflammatory sites. *Science* **325**, 612-616 (2009).
- 17 Darby, W.J., McNutt, K.W., & Todhunter, E.N., Niacin. *Nutr Rev* **33**, 289-297 (1975).
- 18 Attia, E., Wolk, S., Cooper, T., Glasofer, D., & Walsh, B.T., Plasma tryptophan during weight restoration in patients with anorexia nervosa. *Biol Psychiatry* **57**, 674-678 (2005).
- 19 Danilczyk, U. *et al.*, Essential role for collectrin in renal amino acid transport. *Nature* **444**, 1088-1091 (2006).
- 20 Stratigos, J.D. & Katsambas, A., Pellagra: a still existing disease. *Br J Dermatol*

96, 99-106 (1977).

- 21 Nenci, A. *et al.*, Epithelial NEMO links innate immunity to chronic intestinal inflammation. *Nature* **446**, 557-561 (2007).
- 22 Chalaris, A. *et al.*, Critical role of the disintegrin metalloprotease ADAM17 for intestinal inflammation and regeneration in mice. *J Exp Med* **207**, 1617-1624 (2010).
- 23 Ghosh, H.S., McBurney, M., & Robbins, P.D., SIRT1 negatively regulates the mammalian target of rapamycin. *PloS one* **5**, e9199 (2010).
- 24 Broer, A. *et al.*, Impaired nutrient signaling and body weight control in a Na⁺ neutral amino acid cotransporter (Slc6a19)-deficient mouse. *J Biol Chem* **286**, 26638-26651 (2011).
- 25 Farkas, S. *et al.*, Rapamycin decreases leukocyte migration in vivo and effectively reduces experimentally induced chronic colitis. *Int J Colorectal Dis* **21**, 747-753 (2006).
- 26 Reinisch, W. *et al.*, A multicenter, randomized, double-blind trial of everolimus versus azathioprine and placebo to maintain steroid-induced remission in patients with moderate-to-severe active Crohn's disease. *Am J Gastroenterol* **103**, 2284-2292 (2008).
- 27 Mastroianni, J.R. & Ouellette, A.J., Alpha-defensins in enteric innate immunity: functional Paneth cell alpha-defensins in mouse colonic lumen. *J Biol Chem* **284**, 27848-27856 (2009).
- 28 Salzman, N.H. *et al.*, Enteric defensins are essential regulators of intestinal microbial ecology. *Nat Immunol* **11**, 76-83 (2010).
- 29 Weichhart, T., Mammalian target of rapamycin: a signaling kinase for every

aspect of cellular life. *Methods Mol Biol* **821**, 1-14 (2012).

- ³⁰ Abu-Qurshin, R. *et al.*, Crohn's disease associated with pellagra and increased excretion of 5-hydroxyindolacetic acid. *Am J Med Sci* **313**, 111-113 (1997).

Supplementary Information is linked to the online version of the paper at www.nature.com/nature.

Acknowledgements We would like to thank all members of our laboratories for helpful discussions. T.H. was supported by grants from the European Respiratory Society and EuGeneHeart. T.P. is supported by a Marie Curie IIF. J.M.P. is supported by grants from IMBA, the Austrian Ministry of Sciences, the Austrian Academy of Sciences, GEN-AU (AustroMouse), an EU ERC Advanced Grant, and the EU network grants EuGeneHeart, ApoSys, INFLA-Care. P.R. is supported by the BMBF Network “Systematic Genomics of chronic inflammation”, the DFG Cluster of Excellence Inflammation at Interfaces, SPP1399, SFB877, and the DFG project RO1394.

Author Contributions T.H. and T.P. performed most experiments together with J.T., H.I., M.P., V.S., R.H., and T.H.; A.R. and P.R. performed sequencing and analysis of the gut microbiome. S.C., D.S., F.V., and H.C. analyzed Collectrin and Ace2 expression in gut and kidney. A.R. and B.W. performed amino acid analysis in serum. K.K. and A.F. provided essential mouse strains. J.M.P. and P.R. coordinated the project, and together with T.H. and T.P. wrote the manuscript and designed the experiments.

Author Information

Reprints and permissions information is available at www.nature.com/reprints. The authors declare no competing financial interest. Correspondence and requests for materials should be addressed to J.M.P. (josef.penninger@imba.oeaw.ac.at) or P.R. (p.rosenstiel@mucosa.de).

Figure legends

Figure 1. *Ace2* deficiency and protein malnutrition worsen DSS-induced colitis.

a, Colon histopathology, **b**, Percent weight loss and **c**, diarrhea scores in control and DSS-treated *Ace2*^{+/y} and *Ace2*^{-y} littermates. Note crypt damage (arrowheads), ulcerations (arrow), and infiltration of inflammatory cells (asterixes) in DSS-treated *Ace2*^{-y} mice. H&E staining on day 7 after DSS challenge. Scale bars, 100µm. **d**, Colon histopathology (H&E, day 4 after DSS challenge; scale bars, 100µm), **e**, percent weight loss, and **f**, diarrhea scores of DSS-treated *Ace2*^{+/y} and *Ace2*^{-y} littermates fed either normal chow (Control) or a protein free diet (PFD; <0.2% protein). All values are mean ± SEM of 5-9 mice per group. **P*<0.05, ***P*<0.01 comparing DSS-treated *Ace2*^{+/y} vs. *Ace2*^{-y} littermates or *Ace2*^{+/y} mice on normal diet vs. PFD (paired-t-test).

Figure 2. Rescue of severe colitis with nicotinamide or tryptophan di-peptides.

a, Colon histopathology (H&E, day 10 after DSS challenge; scale bars, 100µm), **b**, percent weight loss, and **c**, diarrhea scores of DSS-treated *Ace2*^{+/y} and *Ace2*^{-y} littermates that received vehicle or nicotinamide (NAM) in their drinking water. NAM treatment was started 3 days before DSS challenge. **d**, Colon histopathology (H&E, day 7; scale bars, 100µm), **e**, percent weight loss, and **f**, crypt injury scores of *Ace2*^{+/y} and *Ace2*^{-y} mice fed a di-peptidic tryptophan diet (Trp+) or normal chow (Control). Values are mean ± SEM of 3-10 mice per group. **P*<0.05, ***P*<0.01 comparing *Ace2*^{-y} mice on normal vs. Trp+ diet or vehicle vs. NAM treated *Ace2*^{-y} mice. ### *P*<0.01 comparing *Ace2*^{+/y} vs. *Ace2*^{-y} mice (paired-t-test).

Figure 3. Tryptophan controls antimicrobial peptides and mTOR activity.

a,b, Relative mRNA expression levels of antimicrobial peptides in epithelial cells isolated from the small intestine of **a**, unchallenged *Ace2*^{+/y} and *Ace2*^{-y} littermates, and **b**, *Ace2*^{+/y} mice fed a tryptophan free diet (Trp-) or normal chow (Control). **c**, Relative Defa1 mRNA expression in *Ace2*^{+/y} and *Ace2*^{-y} littermates fed a di-peptidic tryptophan diet (Trp+) or normal chow (Control) for 10 days. **d,e**, Immunohistochemistry to detect levels of phosphorylated S6 (brown) in the small intestine of **d**, unchallenged *Ace2*^{+/y} and *Ace2*^{-y} littermates or **e**, *Ace2*^{-y} mice fed a Trp+ or normal chow diet (Control). Scale bars, 200µm. **f**, Colon histopathology (H&E, day 8; scale bars, 100µm) of DSS treated wild type mice receiving vehicle or RAPA i.p., initiated 6 days before DSS challenge. Values are mean ± SEM of 5-6 mice per group. **P*<0.05, ***P*<0.01 comparing *Ace2*^{+/y} vs. *Ace2*^{-y} mice; #*P*<0.05, ##*P*<0.01 comparing *Ace2*^{+/y} mice on normal vs. tryptophan free diet or control vs. Trp+ diet in *Ace2*^{+/y} or *Ace2*^{-y} mice (paired-t-test).

Figure 4. Altered gut bacteria from *Ace2* mutant mice can confer susceptibility to colitis.

a, Principal coordinate analysis and **b**, Principal coordinate analysis based on unweighted Unifrac analysis showing similarity among ileocaecal bacterial communities in *Ace2*^{+/y} and *Ace2*^{-y} mice fed a di-peptidic tryptophan diet (Trp+) or normal chow (Control) for 10 days. Distances were calculated by Bray-Curtis algorithm from OTU (97% similarity) abundance. Only the two axis with high *R*² values are shown (axis 1, *R*²= 0.335; axis 2, *R*² squared = 0.8116). Each dot represents data from an individual animal. **c**, Comparison of microbial communities in *Ace2*^{+/y} and *Ace2*^{-y} mice fed a Trp+ diet or normal chow (Control). The heat map depicts abundance of the top 25 species level OTUs contributing significantly to the axis shown in the weighted

PcoA plot. **d**, Diarrhea scores and **e**, colon histopathology (H&E, day 7; scale bars, 100µm) of DSS challenged germfree mice that received intestinal microbiota from *Ace2^{+/y}* or *Ace2^{-/y}* littermates. Values are mean \pm SEM of 4-6 mice per group. * $P < 0.05$ (paired-t-test).

Methods

Mice. *Ace2*^{ref.4}, *Agtr1a*^{ref.31}, *Apelin*^{ref.32}, *Apj*^{-/- ref.33}, and *Collectrin*^{ref.19} mutant mice have been described previously. All mutant strains were backcrossed onto a C57BL/6 background more than 6 generations. *Apj*^{-/-} mice were on a mixed C57BL/6 x CBA background. In all experiments, only sex-, age-, and background-matched mice were used as controls. Mice were handled and all experiments performed in accordance with institutional guidelines.

Colitis models. Acute colitis in C57BL/6 background mice was induced with 1,5% (w/v) DSS (molecular mass 36-50 kDa; MP Biologicals) dissolved in drinking water ad libitum for the experimental days 1-7 followed by normal drinking water until the end of the experiment. For mixed C57BL/6 x CBA background mice, colitis was induced by feeding mice 5% (w/v) DSS. The DSS solutions were made fresh every day. TNBS colitis was induced by intrarectal administration of 0.1 ml trinitrobenzene sulphonic acid (TNBS; 5 mg/mouse in 40% ethanol), through a feeding needle (Fine Science Tools) 3.5 cm proximal to the anal verge as described previously³⁴. Body weight, diarrhea, and occult blood (Hemoccult) were determined daily. Diarrhea scores were: 0 = no diarrhea; 1 = mild diarrhea; 2 = severe watery diarrhea; 3 = mild diarrhea with blood; 4 = severe watery diarrhea with blood. Colon length was determined at the end of the experiments. Crypt scores, inflammation scores, and disease activity indexes were determined as previously described¹³. For protein-free diet experiment, normal chow (20,8% protein) and protein-free (<0,2% protein) diets were obtained from Ssniff GmbH, Germany. Mice were fed the protein-free, isocaloric diet for 3 days before start of the 1% (w/v) DSS treatment. Due to the severe diarrhea in PFD fed mice, the experiment had to be stopped for ethical reasons on day 4. For nicotinamide (NAM)

treatment, mice were given NAM (Sigma) in their drinking water (0,4 g/l) from 3 days before the first DSS treatment until the end of the experiment. For rapamycin (RAPA) treatment, rapamycin (2 mg/kg/d; LC laboratories) or vehicle were administered intraperitoneally (i.p.) daily for 6 days with or without NAM treatment. The solvent for rapamycin was 0,2% sodium carboxymethylcellulose, 0,25% polysorbate-80 in water. For tryptophan rescue experiments, standard food was supplemented with glycyl-L-tryptophan (Gly-Trp) hydrate (G0144, TCI Europe) (10mg per gramme of dry food) daily for 10 days followed by 1,5% (w/v) DSS treatment. Tryptophan free diet was purchased from Oriental Yeast (Tokyo, Japan) and was fed for 18 days before DSS challenge. For antibiotic treatment, 4-week-old mice were placed on broad spectrum antibiotics ampicillin (1 g/l) and neomycin (0,5 g/l) in drinking water for 4 weeks followed by 1,5% (w/v) DSS treatment. For microbiota transplantation, cecal and ileum extracts were pooled from *Ace2* mutant mice and ACE2 expressing littermates, suspended in 3ml PBS, and administered (0,1 ml per mouse) immediately to sterile-packed 4-week-old Swiss-Webster germ free mice (Taconic). Transplanted mice were maintained in sterile cages for 2 weeks followed by 5% (w/v) DSS treatment. For ACE2 reconstitution, mice received intraperitoneal injections of recombinant mouse ACE2 protein (0,1 mg/kg) (R&D) daily during 1,5% (w/v) DSS treatment.

Serum amino acid analyses. Heparinised serum was mixed with an equal part of a solution containing $20\mu\text{g mL}^{-1}$ each of norleucine (internal standard for Ala, Gly and Val), norvaline (for Leu, Ile and Pro) and parachloro-phenylalanine (for all other amino acids) and deproteinised by ultrafiltration through a 10 kDa cutoff regenerated cellulose ultrafiltration membrane (Millipore). Amino acids were then purified by cation exchange, converted to their chloroformate derivatives and separated and quantified by

GC/MS in the selected ion monitoring (SIM) mode as previously described⁸. Tryptophan ELISA kit (LDN, BA E-2700) was used to assess Trp serum levels upon Gly-Trp treatment.

Purification of intestinal epithelial cells and mucosal cells. Preparation of small intestinal epithelial cells was performed as described³. Briefly, after washing with PBS, segments of small intestines were placed in 1,5mM EDTA in Hank's balanced salt solution (HBSS) without calcium and magnesium and tumbled for 10 min at 37°C. The mucosa was incubated again with HBSS/EDTA for 10 min at 37°C. The supernatant containing the crypts was collected. The number and viability of the crypt epithelial cells were determined by 0,1% trypan blue exclusion. Purity of the epithelial cell preparation was checked by H&E staining. For mucosa preparations jejunum, ileum, or colon were inverted and mucosa was scraped off with a scalpel.

Quantitative RT-PCR. Total RNA was prepared from isolated small intestinal epithelial cells using the RNeasy Mini Kit (Qiagen) in accordance with the manufacturer's instructions. Total RNA (0,2µg) was subjected to reverse transcription using random hexamers (Roche) and SuperscriptII (Invitrogen) followed by quantitative PCR analysis. The following primers were used:

GAPDH, ' 5 -TCATCAACGGGAAGCCCATCAC-3' (forward) and
5'-AGACTCCACGACATACTCAGCACCG-3' (reverse);

β-actin, ' 5 -GGCTGTATTCCCCTCCATCG-3' (forward) and
5'-CCAGTTGGTAACAATGCCATGT-3' (reverse);

Defa1, ' 5 -TCCTCCTCTCTGCCCTYGTCTTG-3' (forward) and
5'-CTCTTCTCCTGGCTGCTCCTC-3' (reverse);

Defa3, ' 5 -GCTAGGGAGCACTTGTTTGC-3' (forward) and

5'-TTGTTTGAGGAAAGGAGGCA-3' (reverse);

Defa4, ' 5 -CCAGGGGAAGATGACCAGGCTG-3' (forward) and

5'-TGCAGCGACGATTTCTACAAAGGC-3' (reverse);

Defa5, ' 5 -AGGCTGATCCTATCCACAAAACAG-3' (forward) and

5'-TGAAGAGCAGACCCTTCTTGGC-3' (reverse);

Defa-rs1, ' 5 -CACCACCCAAGCTCCAAATACACAG-3' (forward) and

5'-ATCGTGAGGACCAAAAGCAAATGG-3' (reverse);

MMP7, ' 5 -ACCCTGTTCTGCTTTGTGTGTC-3' (forward) and

5'-TCTGAGCCTGTTCCCACTGATG-3' (reverse);

Angiogenin1 (ANG1), 5 -CCATGACGCCAAGCCAAAGG-3' (forward) and

5'-GCTTCCATTCGCTCCACAGATG-3' (reverse);

Angiogenin4 (ANG4), 5 -GCTGGGTCTGGTTGTGATTCC-3' (forward) and

5'-AGGCGAGGTTAGCTTTCTTTCC-3' (reverse);

FABP2, ' 5 -AGCTCGGTGTAACTTTCCCTAC-3' (forward) and

5'-AAACCTCTCGGACAGCAATCAG-3' (reverse);

Lyz1, ' 5 -GCCAAGGTCTACAATCGTTGTGAGTTG-3' (forward) and

5'-CAGTCAGCCAGCTTGACACCACG-3' (reverse);

Lyz2, ' 5 -GGCTGGCTACTATGGAGTCAGCCTG-3' (forward) and

5'-GCATTCACAGCTCTTGGGGTTTTG-3' (reverse);

Pla2g2e, ' 5 -AGGATTCCCCCAAGGATGCCAC-3' (forward) and

5'-CAGCCGTTTCTGACAGGAGTTCTGG -3' (reverse);

Ace, ' 5 -TCATCATCCAGTTCCAGTTCCA-3' (forward) and

5'-CGGTGACGAGCCATTCTGT-3' (reverse);

Ace2, ' 5 -TGGTCTTCTGCCATCCGATT-3' (forward) and

5'-CCATCCACCTCCACTTCTCTAA-3' (reverse);

B⁰AT1, ' 5 -AACGCTCATGTATAGCATCTGG-3' (forward) and

5'-CAGCCACAGTGACCACAAC-3' (reverse);

TLR1, ' 5 -AAGAACTCAGGCGAGCAGAG-3' (forward) and

5'-TCTGGTAAGGTTTGC GTTTGG-3' (reverse);

TLR2, ' 5 -CACTTCCAGGTCTTCAGTCTTC-3' (forward) and

5'-ACTATGATTGCGGACACATCTC-3' (reverse);

TLR3, ' 5 -TTGCGTTGCGAAGTGAAGAA-3' (forward) and

5'-AGACAGATACCTCAGGCTTGG-3' (reverse);

TLR4, ' 5 -CAGAACTTCAGTGGCTGGATT-3' (forward) and

5'-TGTACTAGGTTTCGTCAGATTGG-3' (reverse);

TLR5, ' 5 -GAGTGAGGTCAGTCCTGGAG-3' (forward) and

5'-GAGATGAGGCGTCTGGAGAG-3' (reverse);

TLR6, ' 5 -TACCGTCAGTGCTGGAAATAGA-3' (forward) and

5'-TTCTGTCTTGGCTCATGTTGC-3' (reverse);

TLR7, 5'-AGAACCACTCTATGCCTTCAAG-3' (forward) and

5'-GTCTCTTCCGTGTCCACATC-3' (reverse);

TLR8, ' 5 -CCTTCCTTTGTCTATAGAACATGGA-3' (forward) and

5'-CACAAGGATAGCTTCTGGAATAGT-3' (reverse);

TLR9, ' 5 -AGAATCCTCCATCTCCCAACAT-3' (forward) and

5'-CAGACTTCAGGAACAGCCAAT-3' (reverse);

Western blotting and ELISA. Western blotting was performed with standard protocols using isolated primary small intestinal epithelial cells. Primary antibodies reactive to phosphorylated S6 (2215; Cell Signaling), S6 (2217; Cell Signaling), and β -actin

(Sigma) were used. Blots were washed three times in TBST for 30min, incubated with horseradish peroxidase-conjugated secondary antibodies (1:5000 dilution; Jackson ImmunoResearch) for 20min, washed three times in TBST, and visualised with enhanced chemiluminescence. Small intestine was collected for ELISA assays to detect phosphorylated p70S6K or total p70S6K (DYC896, DYC8962; R&D) and colon was collected to determine angiotensin II levels (EK-002-12; Phoenix). To analyse membrane proteins, total membranes were prepared using mannitol buffer (200mM Mannitol, 80mM HEPES, 41mM KOH, pH 7.5 in the presence of protease inhibitor cocktail, Sigma). After homogenisation of the samples using MagNA Lyser green beads (Roche), total membranes were attained by dual consecutive centrifugation steps (100g and 100'000g) at 4°C. 50 µg of total membrane were used per lane for Western blotting. The following primary antibodies were used: anti-Ace2 and anti-Ace (R&D Systems, 1:1000), anti-B⁰AT1 (Pineda, Berlin, Germany, 1:1000), anti-mouse β-actin (Sigma, St Louis, MO, 1:10000). Antibody binding was detected with Immobilon Western Chemiluminescent HRP or AP substrates (Millipore, Billerica, MA) and chemiluminescence visualised with a DIANA III camera (Raytest, Dietikon, Switzerland).

Histology and immunohistochemistry. For histological analysis, 2-4µm thick sections were stained with haematoxylin and eosin (H&E). Immunohistochemistry was performed using the automated Ventana system. Sections were incubated with antibodies against phosphorylated S6 (Cell Signaling), Ace2 (R&D), Ki67 (Novacastra), and cleaved Caspase 3 (Cell Signaling). For phosphorylated S6 and cleaved Caspase 3 staining, a multimer based detection system (UltraMap) was used. In addition, immunofluorescence was performed on cryosections of the kidney and small intestine

using anti-Ace2 antibodies at 1:100, anti-B⁰AT1 at 1:200, and anti-Collectrin at 1:1000 followed by Alexa488-conjugated anti-goat IgG or Alexa594-labelled anti-rabbit IgG.

DNA extraction, pyrosequencing, sequence analysis

Genomic DNA from terminal ileum was extracted by MO BIO PowerSoil DNA Isolation kit with modifications: solution C1 and 20µl of Proteinase K were added to feces and incubated for 2 hrs at 50°C to enhance the lysis. Extracted DNA was quantified by using the Quant-iT PicoGreen dsDNA Assay Kit (Invitrogen). Variable region V1-V2 of the 16S rRNA gene was amplified using the forward primer 5'-**CTATGCGCCTTGCCAGCCCGCTCAGTC**AGAGTTTGATCCTGGCTCAG-3'.

Sequence in Bold face is 454 Life Sciences primer B, the underlined sequence represents the broadly conserved bacterial primer 27F. A two-base linker sequence ('TC') and a four base key (red) were added. The reverse primer (5'-**CGTATCGCCTCCCTCGCGCCATCAGXXXXXXXXXXCAT**TGCTGCCTCCCGTAGGAGT-3') contained the 454 Life Sciences primer A sequence (bold) a unique 10 base multiplex identifier (MIDs designated as XXXXXXXXXXXX) to tag each PCR product, the broad-range bacterial primer 338R, and a 'CA' linker sequence inserted between the MIDs and the rRNA gene primer. In addition, a four base key (red) was added before the MIDs as described earlier³⁵. Replicate PCR reactions were performed for each sample and the respective negative controls. Each reaction consisted of 1 µl (10 pmol/µl) of each primer, 40 ng of DNA, 1 µl of dNTPs mix (10mM stock), 1.0 units of Phusion Hot Start II High-Fidelity DNA Polymerase, and 10 µl of 5x Phusion HF buffer. The final volume of reaction was adjusted to 50 µl. Amplification was performed using an initial denaturation of 3 min at 98°C followed by 30 cycles, denaturation at 98°C for 10 seconds, annealing at 55°C for 30 seconds and elongation at 72°C for 30 seconds.

Final extension was at 72°C for 10 minutes. Amplified products were run on an agarose gel, specific bands excised and amplicons purified using the Qiagen gel purification kit (Qiagen, Hilden, Germany). Concentrations of the eluted and purified amplicons were measured using the Quant-iT PicoGreen dsDNA Assay Kit (Invitrogen). Equal amounts of PCR products were mixed in a single tube and sequenced using Roche 454 titanium chemistry. To generate negative control templates for the PCR reaction, water was taken as a sample during DNA extraction; in all cases, the PCR reactions from these templates were negative.

Sequence analysis: After pyrosequencing, all sequence reads were screened and filtered for quality and length using PANGEA^{ref.36}. Sequences were trimmed and binned by samples using the specific MIDs as barcode. Sequences not meeting the following quality criteria were excluded from further analysis: no perfect match with MID sequences or specific primers; reads shorter than 200 bases; mean quality score < 25; the presence of any ambiguous base; and/or more than 8 homopolymers. Remaining Sequences were aligned using curated SILVA seed database (bacterial reference alignment) as a template with Needleman-Wunsch algorithm³⁷. Sequences not matching with defined core regions and putative chimeric sequences were detected by using the uchime³⁸ command implemented in Mothur³⁹ and were subsequently removed. Sequences were further confirmed to originate from bacteria using the Ribosomal Data base project (RDP) classifier with 60% bootstrap threshold⁴⁰. For all the downstream analysis, 1000 sequences per samples were randomly sampled to normalize the number of reads distribution. In order to reduce the overestimation of distances between sequences, leading and trailing periods from aligned files were removed. Pair wise distances between sequences were calculated, gaps of any length were treated as single

event or mismatch. Sequences with 97 similarity were clustered using average neighbour algorithm in to species level operational taxonomical units (OTUs)⁴¹. Phylogenetic affiliation of each OTU was performed by using RDP taxonomy and template file. To visualize the microbial composition and structure in related on host genotype or treatment, distance matrices between samples were generated based on weighted (Bray-Curtis similarity) and non-weighted (unweighted UniFrac) algorithms and visualised using principal coordinate analysis (PCoA). OTUs contributing for the shifting of the samples along the two axes were ascertained by calculating the Spearman correlation of each OTU with the two axes in the PCoA^{ref.39}. Similarity percentage analysis (SIMPER)⁴² was performed to determine the driving OTUs imparting significant differences among bacterial community composition among genotype and diet groups. The analysis was performed on Bray Curtis similarity coefficient with the program PAST (Paleontological Statistics, ver. 2.14)⁴³.

Bone marrow transplantation. Eight week old *Ace2* mutant and *Ace2* wild type mice were lethally irradiated with 980 cGy (Gammacell 40). Bone marrow cells were harvested from 8-9 week old donor *Ace2* mutant or *Ace2* wild type mice by flushing the femurs and tibias with DMEM. Recipient mice received 5×10^6 bone marrow cells in 250 μ l DMEM from male donor mice by tail vein injection 6 hrs after irradiation. Six weeks after transplantation, the genotypes of haematopoietic cells were determined by PCR using peripheral blood. The PCR primer set for the mutant *Ace2* allele was as follows: 5'-CCAGCTCATTCCTCCCACTC-3' and 5'-CCGGCTGCTCTTTGAGAGGACA-3'. The PCR primer set for wild-type *Ace2* was: 5'-CTTCATTGGCTCCGTTTCTTAGC-3' and 5'-CCGGCTGCTCTTTGAGAGGACA-3'.

Calorimetry cages. Average food intake was measured using indirect calorimetry cages from TSE Systems over a period of seven days following 48 hrs of acclimatisation. Data were analysed with proprietary Phenomaster Analysis Software (TSE Systems) and Microsoft Excel.

Statistical analyses. All data are shown as mean \pm SEM. Measurements at single time points were analyzed by ANOVA and if significant, further analyzed by a two-tailed t-test. $P < 0,05$ was considered to indicate statistical significance.

- ³¹ Ito, M. *et al.*, Regulation of blood pressure by the type 1A angiotensin II receptor gene. *Proc Natl Acad Sci U S A* **92**, 3521-3525 (1995).
- ³² Kuba, K. *et al.*, Impaired heart contractility in Apelin gene-deficient mice associated with aging and pressure overload. *Circ Res* **101**, e32-42 (2007).
- ³³ Ishida, J. *et al.*, Regulatory roles for APJ, a seven-transmembrane receptor related to angiotensin-type 1 receptor in blood pressure in vivo. *J Biol Chem* **279**, 26274-26279 (2004).
- ³⁴ McCafferty, D.M., Miampamba, M., Sihota, E., Sharkey, K.A., & Kubes, P., Role of inducible nitric oxide synthase in trinitrobenzene sulphonic acid induced colitis in mice. *Gut* **45**, 864-873 (1999).
- ³⁵ Rehman, A. *et al.*, Nod2 is essential for temporal development of intestinal microbial communities. *Gut* **60**, 1354-1362 (2011).
- ³⁶ Giongo, A. *et al.*, PANGEA: pipeline for analysis of next generation amplicons. *ISME J* **4**, 852-861(2010).
- ³⁷ Pruesse, E. *et al.*, SILVA: a comprehensive online resource for quality checked

and aligned ribosomal RNA sequence data compatible with ARB. *Nucleic Acids Res* **35**, 7188-7196 (2007).

³⁸ Edgar, R.C., Haas, B.J., Clemente, J.C., Quince, C., & Knight, R., UCHIME improves sensitivity and speed of chimera detection. *Bioinformatics* **27**, 2194-2200 (2011).

³⁹ Schloss, P.D. *et al.*, Introducing mothur: open-source, platform-independent, community-supported software for describing and comparing microbial communities. *Appl Environ Microbiol* **75**, 7537-7541 (2009).

⁴⁰ Wang, Q., Garrity, G.M., Tiedje, J.M., & Cole, J.R., Naive Bayesian classifier for rapid assignment of rRNA sequences into the new bacterial taxonomy. *Appl Environ Microbiol* **73**, 5261-5267 (2007).

⁴¹ Schloss, P.D. & Handelsman, J., Status of the microbial census. *Microbiol Mol Biol Rev* **68**, 686-691 (2004).

⁴² Clarke, K.R., Nonparametric Multivariate Analyses of Changes in Community Structure. *Aust J Ecol* **18**, 117-143 (1993).

⁴³ Hammer, Ø., Harper, D. A. T. & Ryan, P. D. , Past: paleontological statistics software package for education and data analysis. *Palaeontologia Electronica* **4**, art.4 (2004).